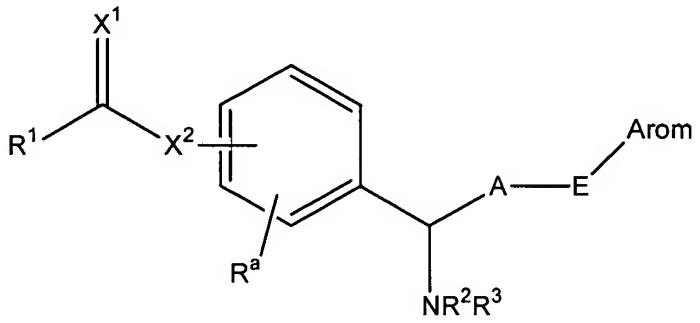
Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

Claims 1 to 45. (canceled)

Claim 46. (currently amended) A compound of formula (I):



(I)

wherein R^1 represents a C_1-C_6 alkyl group, an amino group, a (C_1-C_6 alkyl) amino group, a di(C_1-C_6 alkyl) amino group or a nitrogen-containing saturated heterocyclic group;

R^2 and R^3 are the same or different and represent a hydrogen atom or a C_1-C_6 alkyl group;

Arom represents an unsubstituted phenyl group or a phenyl group substituted at from 1 to 5 positions by one or more substituents which are the same or different and are from the substituent group α;[[;]]

A represents a C_1-C_6 alkylene group;

R^a represents a hydrogen atom, a C_1-C_6 alkyl group or a C_2-C_6 alkenyl group;

E represents an oxygen atom, a sulfur atom or a group of the formula $-NR^4-$, wherein R^4 represents a hydrogen atom or a C_1-C_7 alkanoyl group;

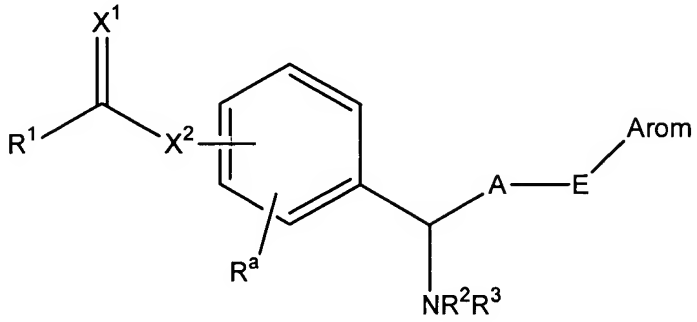
X^1 represents an oxygen atom or a sulfur atom;

X^2 is oxygen and is attached at position C4 of the phenyl ring;

the substituent group α being selected from the group consisting of a halogen atom, C_1-C_6 alkyl group, halogeno C_1-C_6 alkyl group, C_1-C_6 alkoxy group, C_1-C_6 alkylthio group, C_1-C_3 alkylenedioxy group, C_1-C_7 alkanoyl group, C_2-C_7 alkyloxycarbonyl group, amino group, C_1-C_7 alkanoylamino group, hydroxyl group, mercapto group, cyano group, nitro group and carboxyl group;

or a pharmacologically acceptable salt or ester thereof.

Claim 47. (currently amended) A compound of formula (I):



(I)

wherein R^1 represents a C_1-C_6 alkyl group, an amino group, a (C_1-C_6 alkyl) amino group, a di(C_1-C_6 alkyl) amino group or a nitrogen-containing saturated heterocyclic group;

R^2 and R^3 are the same or different and represent a hydrogen atom or a C_1-C_6 alkyl group;

Arom represents an unsubstituted phenyl group or a phenyl group substituted at from 1 to 5 positions by one or more substituents which are the same or different and are from the substituent group α;[[;]]

A represents a C_1-C_6 alkylene group;

R^a represents a hydrogen atom, a C_1-C_6 alkyl group or a C_2-C_6 alkenyl group;

E represents an oxygen atom, a sulfur atom or a group of the formula $-NR^4-$, wherein R^4 represents a hydrogen atom or a C_1-C_7 alkanoyl group;

X^1 represents an oxygen atom or a sulfur atom;

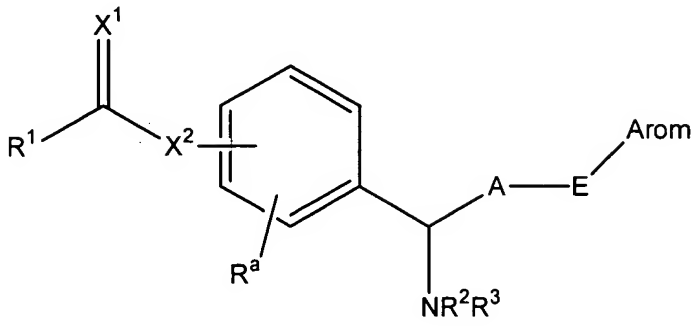
X^2 is oxygen and is attached at position C4 of the phenyl ring;

wherein the group of formula R^1-C(=X^1)- is a (C_1-C_4 alkyl) carbamoyl group or a di(C_1-C_4 alkyl) carbamoyl group;

the substituent group α being selected from the group consisting of a halogen atom, C_1-C_6 alkyl group, halogeno C_1-C_6 alkyl group, C_1-C_6 alkoxy group, C_1-C_6 alkylthio group, C_1-C_3 alkylenedioxy group, C_1-C_7 alkanoyl group, C_2-C_7 alkyloxycarbonyl group, amino group, C_1-C_7 alkanoylamino group, hydroxyl group, mercapto group, cyano group, nitro group and carboxyl group;

or a pharmacologically acceptable salt or ester thereof.

Claim 48. (currently amended) A compound of formula (I):



(I)

wherein R^2 and R^3 are the same or different and represent a hydrogen atom or a C_1-C_6 alkyl group;

Arom represents an unsubstituted phenyl group or a phenyl group substituted at from 1 to 5 positions by one or more substituents which are the same or different and are from the substituent group α;**[[;]]**

A represents a C_1-C_6 alkylene group;

R^a represents a hydrogen atom, a C_1-C_6 alkyl group or a C_2-C_6 alkenyl group;

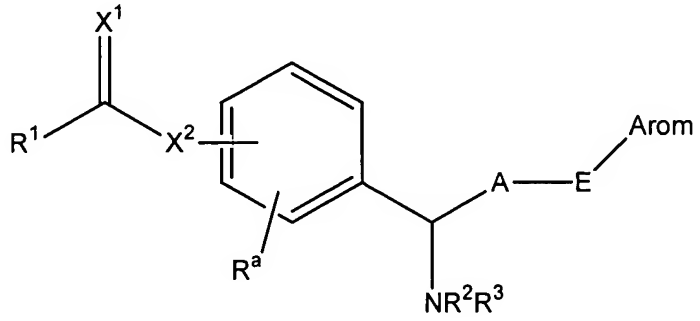
E represents an oxygen atom, a sulfur atom or a group of the formula $-NR^4-$, wherein R^4 represents a hydrogen atom or a C_1-C_7 alkanoyl group;

X^2 is oxygen and is attached at position C4 of the phenyl ring;

wherein the group of formula R^1-C(=X^1)- is a dimethylcarbamoyl group or an ethylmethylcarbamoyl group;

the substituent group α being selected from the group consisting of a halogen atom, C_1-C_6 alkyl group, halogeno C_1-C_6 alkyl group, C_1-C_6 alkoxy group, C_1-C_6 alkylthio group, C_1-C_3 alkylenedioxy group, C_1-C_7 alkanoyl group, C_2-C_7 alkyloxycarbonyl group, amino group, C_1-C_7 alkanoylamino group, hydroxyl group, mercapto group, cyano group, nitro group and carboxyl group; or a pharmacologically acceptable salt or ester thereof.

Claim 49. (currently amended) A compound of formula (I):



(I)

wherein R^1 represents a C_1-C_6 alkyl group, an amino group, a (C_1-C_6 alkyl) amino group, a di(C_1-C_6 alkyl) amino group or a nitrogen-containing saturated heterocyclic group;

R^2 and R^3 are the same or different and represent a hydrogen atom or a C_1-C_6 alkyl group;

Arom is a phenyl group substituted at one or two positions by one or more substituents which are the same or different and are from a substituent group αl, or a phenyl group substituted at three positions by halogen atoms;

A represents a C_1-C_6 alkylene group;

R^a represents a hydrogen atom, a C_1-C_6 alkyl group or a C_2-C_6 alkenyl group;

E represents an oxygen atom, a sulfur atom or a group of the formula $-NR^4-$, wherein R^4 represents a hydrogen atom or a C_1-C_7 alkanoyl group;

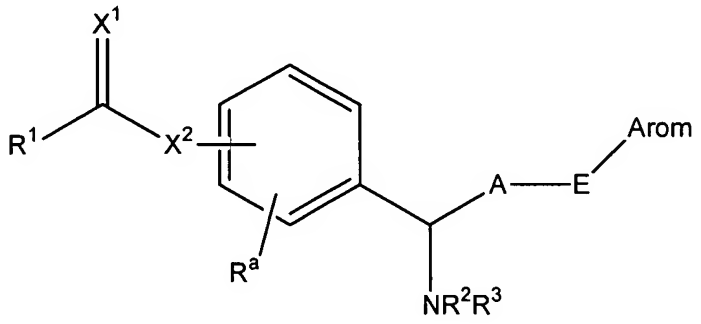
X^1 represents an oxygen atom or a sulfur atom;

X^2 is oxygen and is attached at position C4 of the phenyl ring;

the substituent group αl being selected from the group consisting of a halogen atom, unsubstituted C_1-C_4 alkyl group, C_1-C_4 alkyl group substituted by from 1 to 3 fluorine atoms, C_1-C_4 alkoxy group, C_1-C_4 alkylthio group, methylenedioxy group, ethylenedioxy group, C_1-C_4 alkanoyl group, cyano group and nitro group;

or a pharmacologically acceptable salt ~~or ester~~ thereof.

Claim 50. (currently amended) A compound of formula (I):



(I)

wherein R^1 represents a C_1-C_6 alkyl group, an amino group, a (C_1-C_6 alkyl) amino group, a di(C_1-C_6 alkyl) amino group or a nitrogen-containing saturated heterocyclic group;

R^2 and R^3 are the same or different and represent a hydrogen atom or a C_1-C_6 alkyl group;

Arom is a phenyl group substituted at one or two positions by one or more substituents which are the same or different and are from a substituent group α3, or a phenyl group substituted at three positions by fluorine atoms;

A represents a C_1-C_6 alkylene group;

R^a represents a hydrogen atom, a C_1-C_6 alkyl group or a C_2-C_6 alkenyl group;

E represents an oxygen atom, a sulfur atom or a group of the formula $-NR^4-$, wherein R^4 represents a hydrogen atom or a C_1-C_7 alkanoyl group;

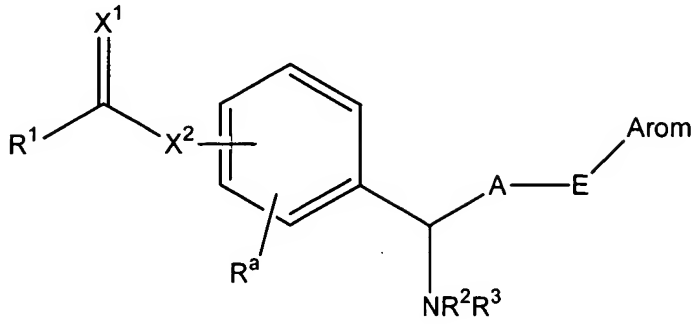
X^1 represents an oxygen atom or a sulfur atom;

X^2 is oxygen and is attached at position C4 of the phenyl ring;

the substituent group α3 being selected from the group consisting of a fluorine atom, chlorine atom, methylthio group, acetyl group, cyano group and nitro group;

or a pharmacologically acceptable salt ~~or ester~~ thereof.

Claim 51. (currently amended) A compound of formula (I):



(I)

wherein R^1 represents a C_1-C_6 alkyl group, an amino group, a (C_1-C_6 alkyl) amino group, a di(C_1-C_6 alkyl) amino group or a nitrogen-containing saturated heterocyclic group;

R^2 and R^3 are the same or different and represent a hydrogen atom or a C_1-C_6 alkyl group;

Arom is a phenyl group substituted at one position by a fluorine atom, a chlorine atomor a nitro group, or a phenyl group substituted at two positions by fluorine atoms;

A represents a C_1-C_6 alkylene group;

R^a represents a hydrogen atom, a C_1-C_6 alkyl group or a C_2-C_6 alkenyl group;

E represents an oxygen atom, a sulfur atom or a group of the formula $-NR^4-$, wherein R^4 represents a hydrogen atom or a C_1-C_7

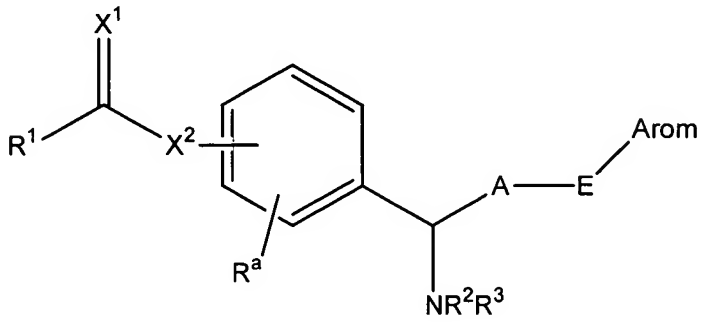
alkanoyl group;

X^1 represents an oxygen atom or a sulfur atom;

X^2 is oxygen and is attached at position C4 of the phenyl ring;

or a pharmacologically acceptable salt ~~or ester~~ thereof.

Claim 52. (currently amended) A compound of formula (I):



(I)

wherein R^1 represents a C_1-C_6 alkyl group, an amino group, a (C_1-C_6 alkyl) amino group, a di(C_1-C_6 alkyl) amino group or a nitrogen-containing saturated heterocyclic group;

R^2 and R^3 are the same or different and represent a hydrogen atom or a C_1-C_6 alkyl group;

Arom is a 4-fluorophenyl group, a 4-chlorophenyl group, a 4-nitrophenyl group or a 3,4-difluorophenyl group;

A represents a C_1-C_6 alkylene group;

R^a represents a hydrogen atom, a C_1-C_6 alkyl group or a C_2-C_6 alkenyl group;

E represents an oxygen atom, a sulfur atom or a group of the formula $-NR^4-$, wherein R^4 represents a hydrogen atom or a C_1-C_7 alkanoyl group;

X^1 represents an oxygen atom or a sulfur atom;

X^2 is oxygen and is attached at position C4 of the phenyl ring;

or a pharmacologically acceptable salt ~~or ester~~ thereof.

Claim 53. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 49, 50, 51 or 52, wherein the group of formula: R^1-C ($=X^1$)- is a carbamoyl group, a (C_1-C_4 alkyl) carbamoyl group, a di(C_1-C_4 alkyl)carbamoyl group, a thiocarbamoyl group, a (C_1-C_4 alkyl) thiocarbamoyl group or a di(C_1-C_4 alkyl) thiocarbamoyl group.

Claim 54. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 49, 50, 51 or 52, wherein the group of formula R^1-C(=X^1)- is a (C_1-C_4 alkyl) carbamoyl group, a di(C_1-C_4 alkyl)carbamoyl group, a (C_1-C_4 alkyl) thiocarbamoyl group or a di(C_1-C_4 alkyl)thiocarbamoyl group.

Claim 55. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 49, 50, 51 or 52, wherein the group of formula R^1-C(=X^1)- is a di(C_1-C_4 alkyl)carbamoyl group.

Claim 56. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 49, 50, 51 or 52, wherein the group of formula R^1-C(=X^1)- is a dimethylcarbamoyl group.

Claim 57. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to

any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein R^3 is a C_1-C_6 alkyl group.

Claim 58. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein R^3 is a methyl group or an ethyl group.

Claim 59. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein R^3 is a methyl group.

Claim 60. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein R^2 is a hydrogen atom or a C_1-C_6 alkyl group.

Claim 61. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein R^2 is a hydrogen atom, a methyl group or an ethyl group.

Claim 62. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein R^2 is a hydrogen atom or a methyl group.

Claim 63. (canceled)

Claim 64. (canceled)

Claim 65. (canceled)

Claim 66. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to

any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein R^a is a hydrogen atom or a methyl group.

Claim 67. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein R^a is a hydrogen atom.

Claim 68. (currently amended) The compound or pharmacologically acceptable salt or ester thereof according to Claims 46 or 47, wherein Arom is a phenyl group substituted at from 1 to 3 positions by one or more substituents which are the same or different and are from the substituent group α, [[,]]

the substituent group α being selected from the group consisting of a halogen atom, C_1-C_6 alkyl group, halogeno C_1-C_6 alkyl group, C_1-C_6 alkoxy group, C_1-C_6 alkylthio group, C_1-C_3 alkylenedioxy group, C_1-C_7 alkanoyl group, C_2-C_7 alkyloxycarbonyl group, amino group, C_1-C_7 alkanoylamino group, hydroxyl group, mercapto group, cyano group, nitro group and carboxyl group.

Claim 69. (canceled)

Claim 70. (previously presented) The compound or pharmacologically acceptable salt thereof according to Claims 46 or 47, wherein Arom is a phenyl group substituted at one or two positions by substituent(s) which are the same or different and are from a substituent group α2, or a phenyl group substituted at three positions by fluorine atoms or chlorine atoms;

substituent group α2 being selected from the group consisting of a fluorine atom, chlorine atom, methyl group, trifluoromethyl group, methoxy group, methylthio group, acetyl group, cyano group and nitro group.

Claim 71. (previously presented) The compound or pharmacologically acceptable salt thereof according to Claims 46 or 47, wherein Arom is a phenyl group substituted at one or two positions by one or more substituents which are the same or different and are from a substituent group α4, or a phenyl group substituted at three positions by fluorine atoms;

substituent group α4 being selected from the group consisting of a fluorine atom, chlorine atom, methylthio group and nitro group.

Claim 72. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein A is a C_1-C_4 alkylene group.

Claim 73. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein A is a methylene group or an ethylene group.

Claim 74. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein A is an ethylene group.

Claim 75. (canceled)

Claim 76. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 47, 48, 49, 50, 51 or 52, wherein E is an oxygen atom.

Claim 77. (canceled)

Claim 78. (canceled)

Claim 79. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 49, 50, 51 or 52, wherein R^1 is an amino group, a (C_1-C_6 alkyl)amino group or a di(C_1-C_6 alkyl)amino group.

Claim 80. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 49, 50, 51 or 52, wherein R^1 is an amino group, a (C_1-C_4 alkyl)amino group or a di(C_1-C_4 alkyl)amino group.

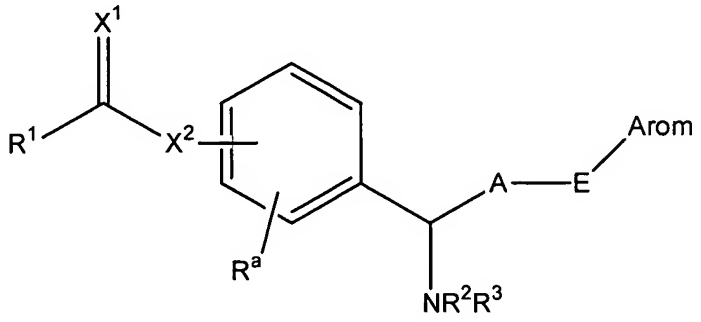
Claim 81. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to

any one of Claims 46, 49, 50, 51 or 52, wherein R^1 is a (C_1-C_4 alkyl)amino group or a di(C_1-C_4 alkyl)amino group.

Claim 82. (previously presented) The compound or pharmacologically acceptable salt or ester thereof according to any one of Claims 46, 49, 50, 51 or 52, wherein X^1 is an oxygen atom.

Claim 83. (currently amended) The compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 46, wherein the compound is 4-[3-(4-nitrophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate.

Claim 84. (previously presented) A compound of the formula (I):



(I)

wherein R^1 represents a C_1-C_6 alkyl group, an amino group, a (C_1-C_6 alkyl)amino group, a di(C_1-C_6 alkyl)amino group or a nitrogen-containing saturated heterocyclic group;

R^2 and R^3 are the same or different and represent a hydrogen atom or a C_1-C_6 alkyl group;

Arom represents an unsubstituted phenyl group or a phenyl group substituted at from 1 to 3 positions by substituents, which are the same or different and are from a substituent group α;

A represents a C_1-C_6 alkylene group;

E represents an oxygen atom, a sulfur atom or a group of the formula $-NR^4-$, wherein R^4 represents a hydrogen atom or a C_1-C_7 alkanoyl group;

X^1 represents an oxygen atom or a sulfur atom;

X^2 is oxygen and is attached at position C4 of the phenyl ring;

the substituent group α being selected from the group consisting of a halogen atom, C_1-C_6 alkyl group, halogeno C_1-C_6 alkyl group, C_1-C_6 alkoxy group, C_1-C_6 alkylthio group, C_1-C_3 alkylenedioxy group, C_1-C_7 alkanoyl group, C_2-C_7 alkyloxycarbonyl

group, amino group, C_1-C_7 alkanoylamino group, hydroxyl group, mercapto group, cyano group, nitro group and carboxyl group;

or a pharmacologically acceptable salt or ester thereof.

Claim 85. (previously presented) A pharmaceutical composition comprising a pharmaceutically effective amount of a compound or pharmacologically acceptable salt or ester thereof according to Claim 46, in combination with a pharmaceutically acceptable carrier.

Claim 86. (previously presented) A pharmaceutical composition comprising a pharmaceutically effective amount of a compound or pharmacologically acceptable salt or ester thereof according to Claim 47, in combination with a pharmaceutically acceptable carrier.

Claim 87. (previously presented) A pharmaceutical composition comprising a pharmaceutically effective amount of a compound or pharmacologically acceptable salt or ester thereof according to

Claim 48, in combination with a pharmaceutically acceptable carrier.

Claim 88. (currently amended) A pharmaceutical composition comprising a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 49, in combination with a pharmaceutically acceptable carrier.

Claim 89. (currently amended) A pharmaceutical composition comprising a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 50, in combination with a pharmaceutically acceptable carrier.

Claim 90. (currently amended) A pharmaceutical composition comprising a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according

to Claim 51, in combination with a pharmaceutically acceptable carrier.

Claim 91. (currently amended) A pharmaceutical composition comprising a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 52, in combination with a pharmaceutically acceptable carrier.

Claim 92. (previously presented) A pharmaceutical composition comprising a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 83, in combination with a pharmaceutically acceptable carrier.

Claims 93 to 101. (canceled)

Claim 102. (previously presented) A method for treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a mammal

comprising administering to a mammal a pharmaceutically effective amount of a compound or a pharmacologically acceptable salt or ester thereof according to Claim 46.

Claim 103. (previously presented) A method for treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a human comprising administering to said human a pharmaceutically effective amount of a compound or pharmacologically acceptable salt or ester thereof according to Claim 46.

Claim 104. (previously presented) A method for treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a human comprising administering to said human a pharmaceutically effective amount of a compound or pharmacologically acceptable salt or ester thereof according to Claim 47.

Claim 105. (previously presented) A method for treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a human comprising administering to said human a pharmaceutically
effective amount of a compound or pharmacologically acceptable salt or ester thereof according to Claim 48.

Claim 106. (currently amended) A method for treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a human comprising administering to said human a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 49.

Claim 107. (currently amended) A method for treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a human comprising administering to said human a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 50.

Claim 108. (currently amended) A method for treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a human comprising administering to said human a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 51.

Claim 109. (currently amended) A method for treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a human comprising administering to said human a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 52.

Claim 110. (canceled)

Claim 111. (previously presented) A method for treating Alzheimer's disease in a human comprising administering to said human a pharmaceutically effective amount of a compound or

pharmacologically acceptable salt or ester thereof according to Claim 46.

Claim 112. (previously presented) A method for treating Alzheimer's disease in a human comprising administering to said human a pharmaceutically effective amount of a compound or pharmacologically acceptable salt or ester thereof according to Claim 47.

Claim 113. (previously presented) A method for treating Alzheimer's disease in a human comprising administering to said human a pharmaceutically effective amount of a compound or pharmacologically acceptable salt or ester thereof according to Claim 48.

Claim 114. (currently amended) A method for treating Alzheimer's disease in a human comprising administering to said human a pharmaceutically effective amount of a compound or

pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 49.

Claim 115. (previously presented) A method for treating Alzheimer's disease in a human comprising administering to said human a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 50.

Claim 116. (currently amended) A method for treating Alzheimer's disease in a human comprising administering to said human a pharmaceutically effective amount of a compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 51.

Claim 117. (currently amended) A method for treating Alzheimer's disease in a human comprising administering to said human a pharmaceutically effective amount of a compound or

pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 52.

Claim 118. (canceled)

Claim 119. (currently amended) The compound or pharmacologically acceptable salt ~~or ester~~ thereof according to Claim 46, wherein R^1 is a dimethylamino group, X^1 and X^2 are both oxygen, Ra is H, R^2 is hydrogen, R^3 is methyl, A is $-C_2H_4$, E is oxygen and Arom is a phenyl group substituted in the 4-position by a NO_2 group.

Claim 120. (currently amended) The compound according to claim 46, wherein the compound is selected from the group consisting of

4-[3-(4-fluorophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate,
4-[3-(3-fluorophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate,
4-[3-(4-chlorophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate,
4-[3-(3-chlorophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate,
4-[3-(4-nitrophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate,

4-[3-(3,4-difluorophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate,
4-[3-(4-chloro-3-fluorophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate,
4-[3-(2-chloro-4-nitrophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate,
4-[1-dimethylamino-3-(4-fluorophenoxy)propyl]phenyl dimethylcarbamate,
4-[1-dimethylamino-3-(3-fluorophenoxy)propyl]phenyl dimethylcarbamate,
4-[3-(4-chlorophenoxy)-1-dimethylaminopropyl]phenyl dimethylcarbamate,
4-[3-(3-chlorophenoxy)-1-dimethylaminopropyl]phenyl dimethylcarbamate,
4-[3-(4-cyanophenoxy)-1-dimethylaminopropyl]phenyl dimethylcarbamate,
4-[1-dimethylamino-3-(4-nitrophenoxy)propyl]phenyl dimethylcarbamate,
4-[3-(3,4-difluorophenoxy)-1-dimethylaminopropyl]phenyl dimethylcarbamate,
4-[3-(2-chloro-4-nitrophenoxy)-1-dimethylaminopropyl]phenyl dimethylcarbamate,
4-[3-(4-nitrophenylsulfanyl)-1-methylaminopropyl]phenyl dimethylcarbamate,
~~4-(1-methylamino-3-p-toluyloxypropyl)phenyl dimethylcarbamate hydrochloride~~[[,]]

4-[1-methylamino-3-[(4-trifluoromethyl)phenoxy]propyl]phenyl dimethylcarbamate,
4-[3-(4-cyanophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate and
4-[1-methylamino-3-(3-nitrophenoxy)propyl]phenyl dimethylcarbamate
or a pharmacologically acceptable salt ~~or ester~~ thereof, or 4-(1-methylamino-3-p-toluyloxypropyl)phenyl dimethylcarbamate hydrochloride.

Claim 121. (currently amended) The compound according to claim 46, wherein the compound is 4-[3-(4-chlorophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate or a pharmacologically acceptable salt ~~or ester~~ thereof.

Claim 122. (currently amended) The compound according to claim 46, wherein the compound is (S)-4-[3-(4-nitrophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate or a pharmacologically acceptable salt ~~or ester~~ thereof.

Claim 123. (currently amended) The compound according to claim 46, wherein the compound is (S)-4-[3-(4-chlorophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate or a pharmacologically acceptable salt ~~or ester~~ thereof.

Claim 124. (previously presented) The compound according to claim 46, wherein the compound is (S)-4-[3-(4-nitrophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate hydrochloride.

Claim 125. (previously presented) The compound according to claim 46, wherein the compound is (S)-4-[3-(4-chlorophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate hydrochloride.

Claim 126. (previously presented) The compound according to claim 46, wherein the compound is (S)-4-[3-(4-nitrophenoxy)-1-methylaminopropyl]phenyl dimethylcarbamate hemifumarate.

Claim 127. (previously presented) The compound according to claim 46, wherein the compound is (S)-4-[3-(4-chlorophenoxy)-1-methylaminopropyl] phenyl dimethylcarbamate hemifumarate.

Claim 128. (currently amended) A method of treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a human comprising administering to said human a pharmaceutically ~~acceptable~~ effective amount of a compound or a pharmaceutically acceptable salt thereof according to any one of claims 83, 121, 122 or 123.

Claim 129. (currently amended) A method of treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a human comprising administering to said human a pharmaceutically ~~acceptable~~ effective amount of a compound according to any one of claims 124, 125, 126 or 127.

Claim 130. (currently amended) A method of treating depression, Huntington's chorea, Pick's disease, tardive dyskinesia, a compulsive disorder or a panic disorder in a human comprising

administering to said human a pharmaceutically ~~acceptable~~ effective of a compound according to claim 126.

Claim 131. (currently amended) A method of treating Alzheimer's disease in a human comprising administering to said human a pharmaceutically ~~acceptable~~ effective amount of a compound or pharmaceutically acceptable salt thereof according to any one of claims 83, 121, 122 or 123.

Claim 132. (currently amended) A method of treating Alzheimer's disease in a human comprising administering to said human a pharmaceutically ~~acceptable~~ effective amount of a compound according to any one of claims 124, 125, 126 or 127.

Claim 133. (currently amended) A method of treating Alzheimer's disease in a human comprising administering to said human a pharmaceutically ~~acceptable~~ effective amount of a compound ~~or pharmaceutically acceptable salt thereof~~ according to claim 126.